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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
09/905,452	07/13/2001	Mohammad Sarwar Nasir	01-660	5761

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EXAMINER

DAVIS, DEBORAH A

ART UNIT	PAPER NUMBER
1641	H

DATE MAILED: 10/08/2002

Please find below and/or attached an Office communication concerning this application or proceeding.

<b>Office Action Summary</b>	<b>Application No.</b>	<b>Applicant(s)</b>
	09/905,452	NASIR ET AL.
<b>Examiner</b>	<b>Art Unit</b>	
Deborah A Davis	1641	

-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --

**Period for Reply**

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If the period for reply specified above is less than thirty (30) days, a reply within the statutory minimum of thirty (30) days will be considered timely.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133).
- Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

## Status

1)  Responsive to communication(s) filed on 25 January 2002 .

2a)  This action is **FINAL**.                            2b)  This action is non-final.

3)  Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

## **Disposition of Claims**

4)  Claim(s) 1-15 is/are pending in the application.

4a) Of the above claim(s) \_\_\_\_\_ is/are withdrawn from consideration.

5)  Claim(s) \_\_\_\_\_ is/are allowed.

6)  Claim(s) 1-15 is/are rejected.

7)  Claim(s) \_\_\_\_\_ is/are objected to.

8)  Claim(s) \_\_\_\_\_ are subject to restriction and/or election requirement.

## Application Papers

9)  The specification is objected to by the Examiner.

10)  The drawing(s) filed on \_\_\_\_\_ is/are: a)  accepted or b)  objected to by the Examiner.

Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).

11)  The proposed drawing correction filed on \_\_\_\_\_ is: a)  approved b)  disapproved by the Examiner.

If approved, corrected drawings are required in reply to this Office action.

12)  The oath or declaration is objected to by the Examiner.

Priority under 35 U.S.C. §§ 119 and 120

13)  Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).

a)  All b)  Some \* c)  None of:

1.  Certified copies of the priority documents have been received.
2.  Certified copies of the priority documents have been received in Application No. \_\_\_\_\_.
3.  Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).

\* See the attached detailed Office action for a list of the certified copies not received.

14)  Acknowledgment is made of a claim for domestic priority under 35 U.S.C. § 119(e) (to a provisional application).

a)  The translation of the foreign language provisional application has been received.

15)  Acknowledgment is made of a claim for domestic priority under 35 U.S.C. §§ 120 and/or 121.

**Attachment(s):**

1)  Notice of References Cited (PTO-892)      4)  Interview Summary (PTO-413) Paper No(s). \_\_\_\_ .  
2)  Notice of Draftsperson's Patent Drawing Review (PTO-948)      5)  Notice of Informal Patent Application (PTO-152)  
3)  Information Disclosure Statement(s) (PTO-1449) Paper No(s)      6)  Other: \_\_\_\_\_

## DETAILED ACTION

### ***Information Disclosure Statement***

1. The listing of references in the specification is not a proper information disclosure statement. 37 CFR 1.98(b) requires a list of all patents, publications, or other information submitted for consideration by the Office, and MPEP § 609 A(1) states, "the list may not be incorporated into the specification but must be submitted in a separate paper." Therefore, unless the references have been cited by the examiner on form PTO-892, they have not been considered.

### ***Drawings***

2. This application has been filed with informal drawings, which are acceptable for examination purposes only. Formal drawings will be required when the application is allowed.

### ***Claim Rejections - 35 USC § 103***

3. The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negatived by the manner in which the invention was made.

4. Claims 1-4 and 8 are rejected under 35 U.S.C. 103(a) as being unpatentable over Nasir et al (Combinatorial Chemistry & High Throughput Screening, 1999, 2, 177-190) in view of Dixon et al (USP#4,835,100).

Nasir et al teaches field tests to determine mycotoxins in human, animal and grain diseases. (pg. 18, last para.). Nasir et al teaches a homogenous assay using fluorescence polarization to analyze these mycotoxins in grains (See abstract). Mycotoxins that are extracted from grains, with a suitable solvent and the sample are added into the antibody solution. A mycotoxin antigen of interest is labeled with a fluorescent molecule (tracer) and is added to the antibody solution. Once the reaction takes place, the fluorescent polarization of the tracer is then measured (pg. 182, para. 1).

Nasir et al does not point out if the particular mycotoxin used was an aflatoxin neither does he make reference to the particular solvent used to extract mycotoxins from a sample.

However, Dixon et al teaches a method and a test kit for detecting an aflatoxin B1 using monoclonal antibodies (See abstract). Dixon et al explains that aflatoxins are toxic metabolites and they can act as potent carcinogens, mutagens and teratogens and are known to occur naturally in wheat and other foods (col. 1, lines 25-34) and (col. 10, lines 45-52). Dixon et al teaches that aflatoxin B1 has to be converted <sup>to</sup> an aflatoxin B1-oxime (aflatoxin B1-carboxymethylamine) when testing for aflatoxins because the aflatoxin B1 lacks the necessary functional groups for conjugation to a label (col. 4, lines 62-68). Dixon et al uses methanol as an extraction solvent (col. 11, lines 36-47).

It would have been obvious to one of ordinary skill in the art to use the method of detecting aflatoxins B1 in food as taught by Dixon et al into the assay of Nasir et al for detecting mycotoxins, to detect toxic levels of contamination in food. It would have

been obvious for Nasir et al to want to detect aflatoxins in grain because certain levels are a public health risk because of the health hazard that they pose to humans and animals. It would have been further obvious to convert aflatoxin B1 into an aflatoxin B1-oxime so that the toxin can be labeled so that fluorescence polarization can be measured with the fluorescent label. This procedure is well known in the art because aflatoxin B1 lacks the necessary functional groups for labeling the toxin. The use of methanol for an extraction solvent is an obvious equivalent of the suitable solvent taught by Nasir et al. With respect to measuring the fluorescence polarization and comparing it with known concentrations of aflatoxin, it would have been obvious to one skilled in the art to do compare toxic levels of aflatoxin in grain to known concentrations in order to determine if said aflatoxins are at high enough levels to pose a health risk.

Claims 5-7 are rejected under 35 U.S.C. 103(a) as being unpatentable over Nasir et al, in view of Dixon et al and in further view of Michel et al (USP#5,741,654).

The teachings of Nasir et al and Dixon et al are set forth above and differ from the instant claims in not particularly pointing out a particular type of fluorescein used in the assay.

However, Michel et al discloses a Fluorescence Polarization assay for the quantification of antibodies in which a variety of fluoresceins are used as detectable moiety components of tracers, such as one mentioned in particular, the 6-aminofluorescein moiety (isomer II of fluorescein) which is one of the preferred moieties of choice in the said assay (col. 8, lines 1-22).

It would have been obvious to one of ordinary skill in the art to employ a fluoresceinamine or its isomers as binding moieties because such structures are well known in the art to work well in Fluorescence Polarization Immunoassays for quantitation of a sample. In addition, the fluorescein used for labeling in this assay would have been a functional equivalent of the fluorescent molecule used for labeling in the assay of Nasir et al - wherein both would have worked equally as well.

5. Claims 9-10 are rejected under 35 U.S.C. 103(a) as being unpatentable over Nasir et al in view of Dixon et al, and further in view of McMahon et al (USP#5,166,078).

The teachings of Nasir et al and Dixon et al are set forth above and differ from the instant claims in not teaching the construction of a standard curve using a plurality of different known concentrations of aflatoxin.

However, McMahon et al teaches a method for measuring a hapten that is poorly soluble in an aqueous solution such as aflatoxins (col. 2, lines 45-53). The invention permits fast, safe, and convenient measurements of haptens, which are either insoluble or unstable in aqueous solution by providing standards that are soluble and stable in aqueous solution. The standards are used to determine the amount of haptens that are present in the assay (col. 1, lines 43-48). To determine the amount of hapten in a sample, the reaction of the hapten and the antibody is compared to the reaction of the hapten-conjugate and the antibody. The conjugates of the invention are used as controls in standard immunoassay (col. 2, lines 29-40). The reactivity of the conjugate

was compared to aflatoxin standards and a standard curve was created relating aflatoxin levels to aflatoxin-conjugate levels (col. 3, lines 9-16).

It would have been obvious to one of ordinary skill in the art to use a plurality of aflatoxins in standard solutions having different known concentrations and comparing them with aflatoxin-conjugates to create a standard curve to permit fast, safe and convenient measurements of haptens. Further, one skilled in the art would know that certain levels of aflatoxins found in different amounts of grain are toxic to human and animals and a standard curve is needed to compare those levels that would be of concern.

6. Claims 11-18 are rejected under 35 U.S.C. 103(a) as being unpatentable over Nasir et al in view of Dixon et al (USP#4,835,100).

The teachings of Nasir et al are set forth above and differ from the instant claims in not teaching the assay in the form of a kit.

Dixon et al however discloses a kit for aflatoxins and explains that obvious variations of preparing a kit for convenience will be apparent to those skilled in the art and points out that kits are well developed in the patent arts and literature (col. 12, lines 28-33).

It would have been *prima facie* obvious to one of ordinary skill in the art to take the assay for aflatoxins as taught by Dixon et al, combined with the teachings of Nasir et al, for the determination of Mycotoxins and formulate a kit. Further, it would be convenient to do so because one can enhance sensitivity of a method by providing

reagents as a kit. In addition, the reagents in a kit are available in premeasured amounts, which eliminates the variability that can occur when performing the assay.

***Conclusion***

7. The prior art made of record and not relied upon is considered pertinent to applicant's disclosure.

- A. Peter J. Cotty (USP#5,171,686) provides methods and composition for the control or prevention of aflatoxin contamination of agricultural commodities.
- B. Hart et al (USP#4,772,551) provides a method and a test kit for detecting a trichothecene using monoclonal antibodies.

No claims are allowed.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Deborah A Davis whose telephone number is (703) 308-4427. The examiner can normally be reached on 8-5 Monday thru Friday.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Long Le can be reached on (703) 305-3399. The fax phone numbers for the organization where this application or proceeding is assigned are (703) 308-4242 for regular communications and (703) 308-4242 for After Final communications.

Art Unit: 1641

Any inquiry of a general nature or relating to the status of this application or proceeding should be directed to the receptionist whose telephone number is (703) 308-1123.



Deborah A. Davis  
CM1, 7D16  
September 23, 2002



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